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Substitute for form 1449A/PTO SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT BY APPLICANT <i>(use as many sheets as necessary)</i>		Complete if Known <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%;">Application Number</td> <td>09/834,596</td> </tr> <tr> <td>Filing Date</td> <td>April 13, 2001</td> </tr> <tr> <td>First Named Inventor</td> <td>Watanabe <i>et al.</i></td> </tr> <tr> <td>Group Art Unit</td> <td>1623</td> </tr> <tr> <td>Examiner Name</td> <td>Howard V. Owens, Jr.</td> </tr> <tr> <td>Attorney Docket Number</td> <td>08841.105037 PHAR 2020</td> </tr> </table>		Application Number	09/834,596	Filing Date	April 13, 2001	First Named Inventor	Watanabe <i>et al.</i>	Group Art Unit	1623	Examiner Name	Howard V. Owens, Jr.	Attorney Docket Number	08841.105037 PHAR 2020
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Attorney Docket Number	08841.105037 PHAR 2020														
Sheet	1	of	7												

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U.S. PATENT DOCUMENTS

Examiner Initials *	Cite No. ¹	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages/Relevant Figures Appear
		Number	Kind Code (if known)			
<i>Ho</i>	AA	3,480,613		Walton <i>et al.</i>	11-25-1969	
	AB	5,977,061		Holy <i>et al.</i>	11-02-1999	
	AC	6,340,690	B1	Bachand <i>et al.</i> (Idenix Pharm.)	01-22-2002	
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	AF	6,573,248	B2	Ramasamy <i>et al.</i>	06-03-2003	
	AG	2002/0055483	A1	Watanabe <i>et al.</i>	05-09-2002	
	AH	2002/0147160	A1	Bhat <i>et al.</i>	10-10-2002	
	AI	2003/0008841	A1	Devos <i>et al.</i>	01-09-2003	
	AJ	2003/0028013	A1	Wang <i>et al.</i>	02-06-2003	
	AK	2003/0050229	A1	Sommadosi <i>et al.</i> (Idenix Pharm.)	03-13-2003	
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<i>Ho</i>	AM	2003/0087873	A1	Stuyver <i>et al.</i>	05-08-2003	

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		Office ³	Number	Kind Code ² (if known)				
	AN	FR	1,521,076		Merck & Co. Inc.	04-12-1968		
	AO	FR	1,581,628		Merck & Co. Inc.	09-19-1969		
	AP	FR	2,662,165	A1	Univ Pierre et Marie Curie, Paris	11-22-1991		
<i>Ho</i>	AQ	GB	1,163,103	A	Merck & Co. Inc.	09-04-1969		
<i>Ho</i>	AR	GB	1,209,654	A	Merck & Co. Inc.	10-21-1970		
	AS	JP	63-215694	A	Yamasa Shoyu Co. Ltd.	09-08-1988		
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<i>Ho</i>	AV	WO	99/43691	A1	Emory U./Georgia Res. Found.	09-02-1999		

Examiner Signature		Date Considered	6/30/04
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Sheet **2** of **7**

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		Office ³	Number	Kind Code ² (if known)				
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	BB	WO	01/16671	A1	Novirio (Idenix Pharmaceuticals)	03-08-2001		
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	BK	WO	02/18404	A2	F. Hoffmann-La Roche AG	03-07-2002		
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	BN	WO	02/057287	A2	Merck & Co. Inc.; Isis Pharm.	07-25-2002		
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X/HO	BZ	WO	03/062257	A1	Ribapharm Inc.	07-31-2003		

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Howard V. Owens, Jr.

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		Office ³	Number	Kind Code ² (if known)				
<i>HO</i>	CA	WO	03/063771	A2	Pharmasset Ltd.	08-07-2003		
<i>HO</i>	CB	WO	03/068162	A2	Pharmasset Ltd.	08-21-2003		
<i>HO</i>	CC	WO	03/072757	A2	Biota Inc.	09-04-2003		
<i>HO</i>	CD	WO	03/093290	A2	Genelabs Technologies Inc.	11-13-2003		

OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	1 st
	CE	ALTMANN, K.H., <i>et al.</i> , "The Synthesis of 1'-methyl carboacyclic thymidine and its effect on nucleic acid duplex stability", <i>Synlett</i> , Thieme Verlag, Stuttgart, Gc, October 1994, 10, 853-855	
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Harold Cas

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Sheet 4 of 7

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OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

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	DA	EARKAS, J., "Nucleic acids components and their analogues. LXXIX. Synthesis of methyl 1-deoxy-D-psicofuranosides substituted at C(1) with halo atoms or a mercapto group," <i>Collect. Czech. Chem. Commun.</i> 1966, 31, 1535-1543.	
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	DE	HARAGUCHI, K., <i>et al.</i> , "Preparation and reactions of 2'- and 3'-vinyl bromides of uracil-nucleosides: versatile synthons for anti-HIV agents," <i>Tetrahedron Letters</i> , 1991, 32(28), 3391-3394.	
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	DN	JOHNSON, C.R., <i>et al.</i> , "3'-C-trifluoromethyl ribonucleosides, <i>Nucleosides & Nucleotides</i> , 1995, 14, 185-194.	

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Sheet **5** of **7**

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	EI	MIKHAILOV, S.N., et al., "Synthesis and properties of 3'-C-methylnucleosides and their phosphoric esters," <i>Carbohydrate Research</i>, 1983, 124, 75-96.	
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	EK	MIKHAILOV, S.N., et al., "Substrate properties of C'-methyl nucleoside and C'-methyl-2'-deoxynucleoside 5'-triphosphates in RNA and DNA synthesis reactions catalysed by RNA and DNA polymerases," <i>Nucleosides & Nucleotides</i>, 1991, 10, 339-343.	
	EL	NUTT, R.F., et al., "Branched-chain sugar nucleosides. III. 3'-C-methyladenine," <i>J. Org. Chem.</i> 1968, 33, 1789-1795.	

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Sheet 6 of 7

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Application Number	09/834,596
Filing Date	April 13, 2001
First Named Inventor	Watanabe <i>et al.</i>
Group Art Unit	1623
Examiner Name	Howard V. Owens, Jr.
Attorney Docket Number	08841.105037 PHAR 2020

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OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ⁴
	FA	OIVANEN, M., <i>et al.</i> , "Additional evidence for the exceptional mechanism of the acid-catalysed hydrolysis of 4-oxypyrimidine nucleosides: hydrolysis of 1-(1-alkoxyalkyl)uracils. Seconucleosides. 3'-C-alkyl nucleosides and nucleosides 3',5'-cyclic monophosphates," <i>J. Chem. Soc. Perkin Trans.</i> , 1994, 2, 309-314.	
	FB	ONG, S.P., <i>et al.</i> , "Synthesis of 3'-C-methyl adenosine and 3'-C-methyluridine diphosphates and their interaction with the ribonucleoside diphosphate reductase from <i>Corynebacterium nephridii</i> ," <i>Biochemistry</i> , 1992, 31, 11210-11215.	
	FC	PAN-ZHOU X-R., <i>et al.</i> , "Differential effects of antiretroviral nucleoside analogs on mitochondrial function in HepG2 cells," <i>Antimicrob Agents Chemother</i> 2000, 44(no.3), 496-503.	
	FD	ROSENTHAL, A., <i>et al.</i> , "Branched-chain sugar nucleosides. Synthesis of 3'-C-ethyl (and 3'-C-butyl)uridine," <i>Carbohydrate Research</i> , 1980, 79, 235-242.	
	FE	SAMANO, V., <i>et al.</i> , "Nucleic acid related compounds. 77. 2',3'-didehydro-2',3'-dideoxy-2'-(and 3')-methyl nucleosides via [3,3]-sigmatropic rearrangements of 2'-(and 3')-methylene-3'-(and 2')-O-thiocarbonyl derivatives and radical reduction of a 2'-chloro-3'-methylene analogue," <i>Can. J. Chem.</i> , 1993, 71, 186-191.	
	FF	SAMANO, V., <i>et al.</i> , "Synthesis and radical-induced ring-opening reactions of 2'-deoxyadenosine-2'-spirocyclopropane and its uridine analogs. Mechanistic probes for ribonucleotide reductases," <i>J. Am. Chem. Soc.</i> , 1992, 114, 4007-4008.	
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	FJ	SOMMADOSSI J-P., <i>et al.</i> , "Comparison of cytotoxicity of the (-) and (+)-enantiomer of 2',3'-dideoxy-3'-thiaeytidine in normal human bone marrow progenitor cells," <i>Biochemical Pharmacology</i> , 1992, 44:1921-1925.	
	FK	SOMMADOSSI J-P., <i>et al.</i> , "Toxicity of 3'-azido-3'-deoxythymidine and 9-(1,3-dihydroxy-2-propoxymethyl)guanine for normal human hematopoietic progenitor cells in vitro," <i>Antimicrobial Agents and Chemotherapy</i> , 1987, 31(No. 3), 452-454.	
	FL	TRITSCH, D., <i>et al.</i> , "3'-β-ethynyl and 2'-deoxy-3'-β-ethynyl adenosines: first 3'-β-branched-adenosines substrates of adenosine deaminase," <i>Biorganic & Medicinal Chemistry Letters</i> , 2000, 10, 139-141.	

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Sheet	7	of	7
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Application Number	09/834,596
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First Named Inventor	Watanabe <i>et al.</i>
Group Art Unit	1623
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Attorney Docket Number	08841.105037 PHAR 2020

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Sheet 1 of 3

Application Number	09/834,596
Filing Date	13 April 2001
First Named Inventor	Watanabe, K.
Group Art Unit	1614
Examiner Name	Not Assigned
Attorney Docket Number	PHARM1

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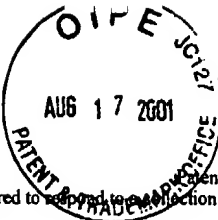
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INFORMATION DISCLOSURE STATEMENT BY APPLICANT (use as many sheets as necessary)		Filing Date	04/13/01
		First Named Inventor	WATANABE
		Group Art Unit	1614
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Sheet	2 of 3		

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Ho		Fedorov, I., et al., XIII International Round Table: Nucleosides, Nucleotides and Their Biological Applications. Montpellier, France, September 6-10, 1998. Poster 35.	
Ho		Kato, K., et al., XIII International Round Table: Nucleosides, Nucleotides and Their Biological Applications. Montpellier, France, September 6-10, 1998. Poster 49.	
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INFORMATION DISCLOSURE STATEMENT BY APPLICANT (use as many sheets as necessary)		Filing Date	04/13/01
		First Named Inventor	WATANABE
		Group Art Unit	1614
		Examiner Name	Not Assigned
		Attorney Docket Number	PHARM1
Sheet	3	of	3

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AO		Sekiyama, T., J. Med. Chem., 1998, 41:1284-98	

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